Drug Enforcement Administration

[Docket No. DEA-392]

Importer of Controlled Substances Application: AMRI Rensselaer, Inc.

ACTION: Notice of application.

DATES: Registered bulk manufacturers of the affected basic classes, and applicants therefore, may file written comments on or objections to the issuance of the proposed registration in accordance with 21 CFR 1301.34(a) on or before [INSERT 30 DAYS AFTER PUBLICATION IN THE FEDERAL REGISTER]. Such persons may also file a written request for a hearing on the application pursuant to 21 CFR 1301.43 on or before [INSERT 30 DAYS AFTER PUBLICATION IN THE FEDERAL REGISTER]. **ADDRESS:** Written comments should be sent to: Drug Enforcement Administration, Attention: DEA Federal Register Representative/DRW, 8701 Morrissette Drive, Springfield, Virginia 22152. All requests for hearing must be sent to: Drug Enforcement Administration, Attn: Administrator, 8701 Morrissette Drive, Springfield, Virginia 22152. All requests for hearing should also be sent to: (1) Drug Enforcement Administration, Attn: Hearing Clerk/LJ, 8701 Morrissette Drive, Springfield, Virginia 22152; and (2) Drug Enforcement Administration, Attn: DEA Federal Register Representative/DRW, 8701 Morrissette Drive, Springfield, Virginia 22152. Comments and requests for hearings on applications to import narcotic raw material are not appropriate. 72 FR 3417 (January 25, 2007).

SUPPLEMENTARY INFORMATION:

The Attorney General has delegated his authority under the Controlled Substances Act

to the Administrator of the Drug Enforcement Administration (DEA), 28 CFR 0.100(b).

Authority to exercise all necessary functions with respect to the promulgation and

implementation of 21 CFR part 1301, incident to the registration of manufacturers,

distributors, dispensers, importers, and exporters of controlled substances (other than

final orders in connection with suspension, denial, or revocation of registration) has been

redelegated to the Assistant Administrator of the DEA Diversion Control Division

("Assistant Administrator") pursuant to section 7 of 28 CFR part 0, appendix to subpart

R.

In accordance with 21 CFR 1301.34(a), this is notice that on June 27, 2016, AMRI

Rensselaer, Inc., 33 Riverside Avenue, Rensselaer, New York 12144 applied to be

registered as an importer of poppy straw concentrate (9670), a basic class of controlled

substance listed in schedule II.

The company plans to import the listed controlled substance to manufacture bulk

controlled substance for distribution to its customers.

Dated: July 20, 2017

Demetra Ashley,

Acting Assistant Administrator.

[FR Doc. 2017-15689 Filed: 7/25/2017 8:45 am; Publication Date: 7/26/2017]

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